

ОВ	
CDC Use Only	

REPORT OF OUTBREAK OF SUSPECTED VIRAL GASTROENTERITIS

Viral Gastroenteritis Section DASH Unit 75	Telephone (404) 639-3577 or (404) 639-3607
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Primary contact for epidemiologic investigation	Date
Name	
Agency	
Address	
Outbreak Information	
State Outbreak ID EFORS code Date hea	alth department notified
	mm/dd/yyyy
Date of first case Date of last ca	ase mm/dd/yyyy
Location(s) of outbreak: StateCityCounty (list i	if several)
If multistate, list other states	
Suspected mode of transmission (can check more than one)	
□ Person-to-person □ Foodborne	□ Waterborne
□ Unknown □ Other	
If food or waterborne Implicated food or water source Foodhandler im (can check more	Yes: epidemiologic evidence Yes: laboratory evidence Yes: suspected, but no evidence No
<u>Setting</u> (if there is an additional setting, please add below in comments)	
Nursing home Assisted Living Restaurant / Deli / Cafeteria	Hotel School Daycare Camp
Community Prison Ship Religious Facility Hospit	al Private event Catered Event
Work Place Private Home Other Date	of event (if applicable)
Illness Characteristics	ппи за уууу
Number of persons exposed Number of persons ill Number visiting health care provider Number (Categories NOT mutually exclusive)	er hospitalizedNumber deaths
Symptoms: Number of persons with information Median	incubation period (hours) range
Symptoms: Number of persons with information Median No. with abdominal cramps No. with fever Median No. with diarrhea No. with vomiting Median	· · · · · · · · · · · · · · · · · · ·
No. with diarrhea No. with vomiting Median	n duration of illness (hours) range
No. with other symptom(s)	
Comments:	

Part II

Facsim	ile	E-Mail	
		Number of vomitus specimens:	submitted
Yes	No	Results (if known)	
Yes	No	Results (if known)	
lected fro	m:	ill persons control perso	ons
of convale	escent ser	a	
34 (DASH Formula) 3 weeks aft	orm). Acute ter the collec	be stored and shipped frozen in plastic sera should be collected within 7 days ction of acute sera.	of onset of symptoms and
No	Туре	Date Colle	ected
			mm/dd/yyyy
mm	/dd/yyyy	Specimen type	
mm	/dd/yyyy	Specimen type	
, etc):			
are and e	necimen	tyne for samples submitted to	o CDC for testing
pecimen ⁻	Гуре'	Date of Collection (mm/dd/yyy	yy) Additional Information ²
	Yes Yes d from ill per on frozen re lected from of convale blogic specim 34 (DASH F 1 3 weeks aff No mm mm , etc): ers and s	Yes No Yes No Yes No d from ill persons should on frozen refrigerant pa lected from: of convalescent ser blogic specimens should 34 (DASH Form). Acute 1 3 weeks after the collect No Type mm/dd/yyyy mm/dd/yyyy , etc):	Number of vomitus specimens

¹e.g. Bulk stool, vomitus, serum, rectal swab, environmental specimen ²e.g. Foodhandler / ship crew

RECOMMENDATIONS REGARDING SPECIMEN COLLECTION FOR DIAGNOSIS OF NLVs* Clinical Specimens

Stool

Timing. Specimen collection for viral testing should begin on day 1 of the epidemiologic investigation. Any delays to await testing results for bacterial or parasitic agents could preclude establishing a viral diagnosis. Ideally, specimens should be obtained during the acute phase of illness (i.e., within 48--72 hours after onset) while the stools are still liquid or semisolid because the level of viral excretion is greatest then. With the development of sensitive molecular assays, the ability to detect viruses in specimens collected later in the illness has been improved. In specific cases, specimens might be collected later during the illness (i.e., 7--10 days after onset), if the testing is necessary for either determining the etiology of the outbreak or for epidemiologic purposes (e.g., a specimen obtained from an ill foodhandler who might be the source of infection). If specimens are collected late in the illness, the utility of viral diagnosis and interpretation of the results should be discussed with laboratory personnel before tests are conducted.

Number and Quantity. Ideally, specimens from ≥10 ill persons should be obtained during the acute phase of illness. Bulk samples (i.e., 10--50 ml of stool placed in a stool cup or urine container) are preferred, as are acute diarrhea specimens that are loose enough to assume the shape of their containers. Serial specimens from persons with acute, frequent, high-volume diarrhea are useful as reference material for the development of assays. The smaller the specimen and the more formed the stool, the lower the diagnostic yield. Rectal swabs are of limited or no value because they contain insufficient quantity of nucleic acid for amplification. Storage and Transport. Because freezing can destroy the characteristic viral morphology that permits a diagnosis by EM, specimens should be kept refrigerated at 4 C. At this temperature, specimens can be stored without compromising diagnostic yield for 2--3 weeks, during which time testing for other pathogens can be completed. If the specimens have to be transported to a laboratory for testing, they should be bagged and sealed and kept on ice or frozen refrigerant packs in an insulated, waterproof container. If facilities for testing specimens within 2--3 weeks are not available, specimens can be frozen for antigen or PCR testing.

Vomitus

Vomiting is the predominant symptom among children, and specimens of vomitus can be collected to supplement the diagnostic yield from stool specimens during an investigation. Recommendations for collection, storage, and shipment of vomitus specimens are the same as those for stool specimens.

Serum

Timing. If feasible, acute- and convalescent-phase serum specimens should be obtained to test for a diagnostic ≥4-fold rise in IgG titer to NLVs. Acute-phase specimens should be obtained during the first 5 days of symptoms, and the convalescent-phase specimen should be collected from the third to sixth week after resolution of symptoms.

Number and Quantity. Ideally, 10 pairs of specimens from ill persons (i.e., the same persons submitting stool specimens) and 10 pairs from well persons (controls) should be obtained. Adults should provide 5--7 ml of blood, and children should provide 3--4 ml.

Storage. Specimens should be collected in tubes containing no anticoagulant, and the sera should be spun off and frozen. If a centrifuge is not available, a clot should be allowed to form, and the serum should be decanted and frozen. If this step cannot be accomplished, the whole blood should be refrigerated but not frozen.

Environmental Specimens

NLVs cannot be detected routinely in water, food, or environmental specimens. Nevertheless, during recent outbreaks (33--36), NLVs have been detected successfully in vehicles epidemiologically implicated as the source of infection. If a food or water item is strongly suspected as the source of an outbreak, then a sample should be obtained as early as possible and stored at 4 C. If the epidemiologic investigation confirms the link, a laboratory with the capacity to test these specimens should be contacted for further testing. If drinking water is suspected, special filtration (45) of large volumes (i.e., 5--100 liters) of water can concentrate virus to facilitate its detection.